

U.S. PATENT DOCUMENTS

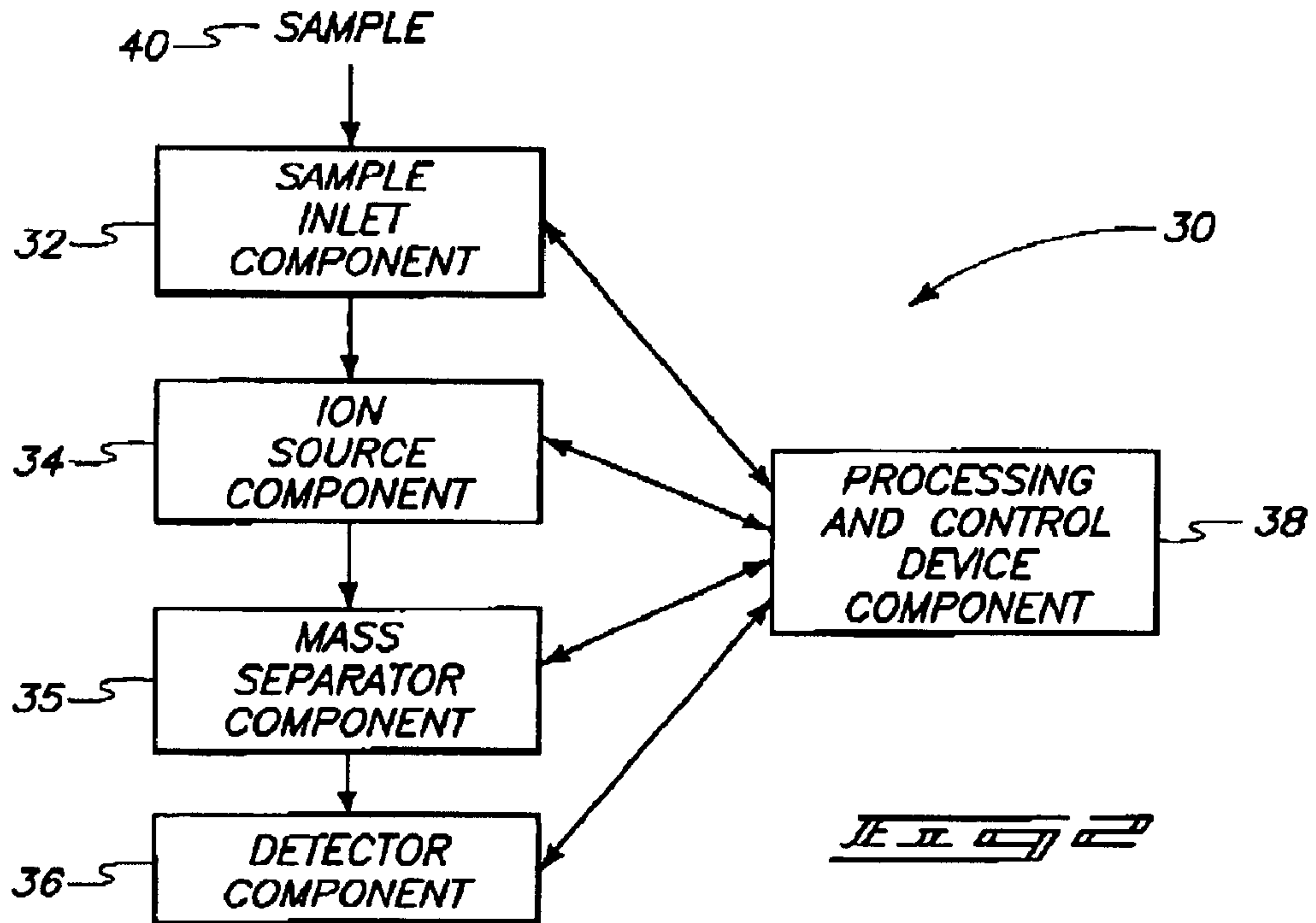
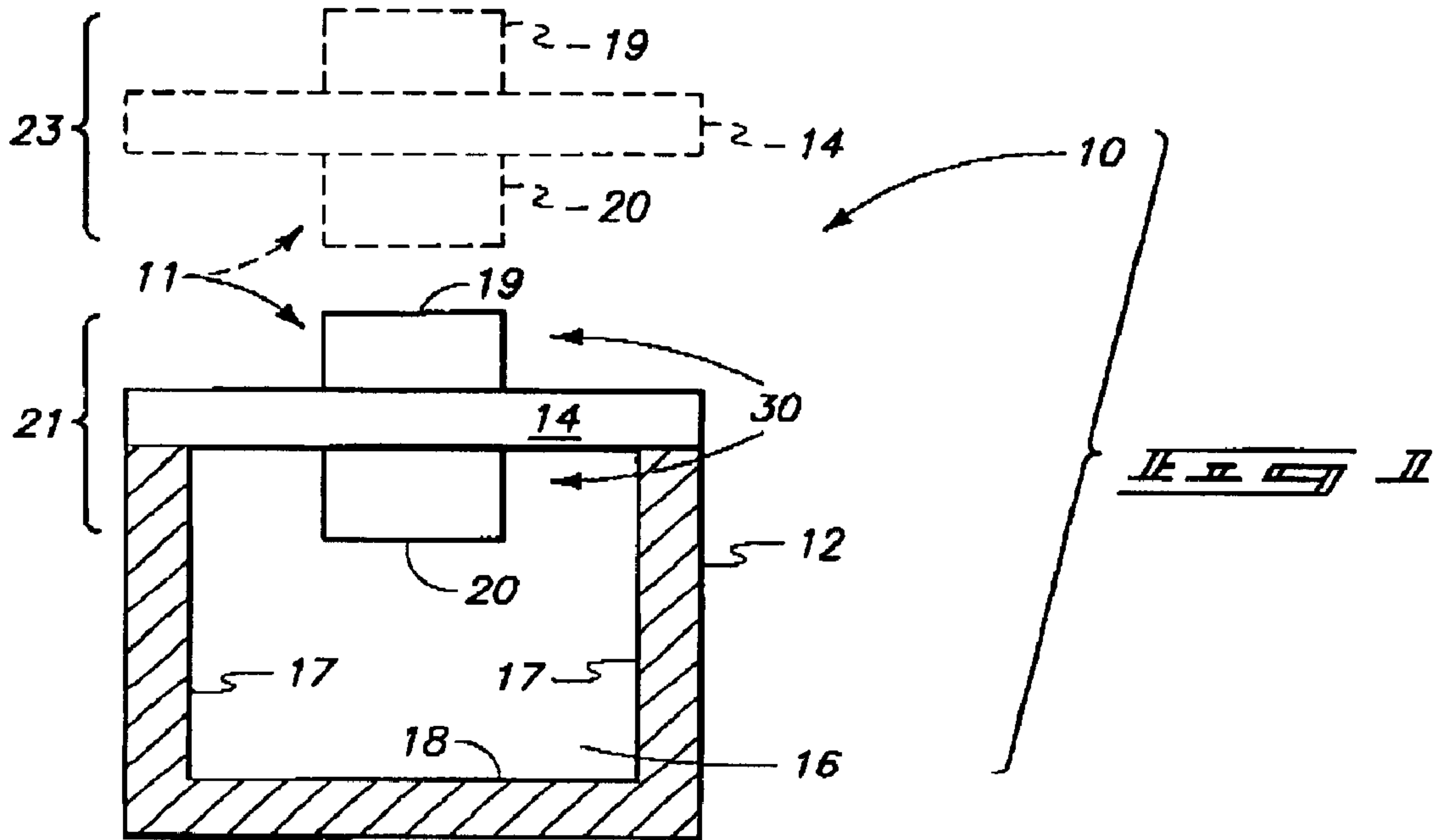
4,755,685 A * 7/1988 Kawanami et al. 250/492.2
 4,849,628 A 7/1989 McLuckey et al.
 5,313,061 A * 5/1994 Drew et al. 250/281
 5,818,055 A * 10/1998 Franzen 250/292
 5,837,883 A * 11/1998 Itoi 73/23.37
 5,844,237 A * 12/1998 Whitehouse et al. 250/288
 5,896,196 A 4/1999 Pinnaduwege
 6,239,429 B1 * 5/2001 Blessing et al. 250/292
 6,329,654 B1 * 12/2001 Gulcicek et al. 250/293
 6,410,914 B1 6/2002 Park et al.
 6,541,768 B2 * 4/2003 Andrien et al. 250/288
 6,593,568 B1 * 7/2003 Whitehouse et al. 250/288
 6,825,466 B2 * 11/2004 Mordekhay 250/288
 2002/0053636 A1 * 5/2002 Shiokawa et al. 250/281

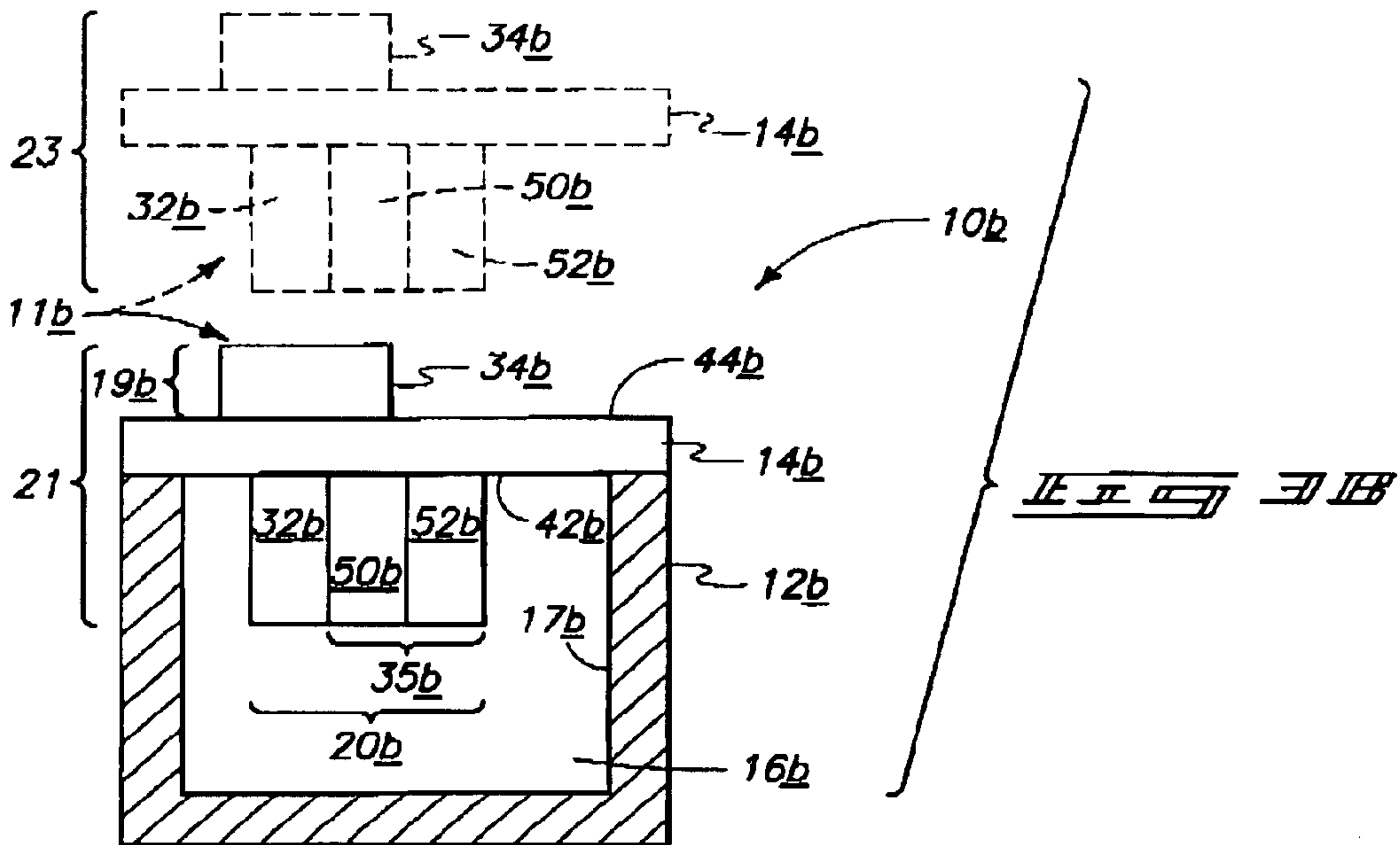
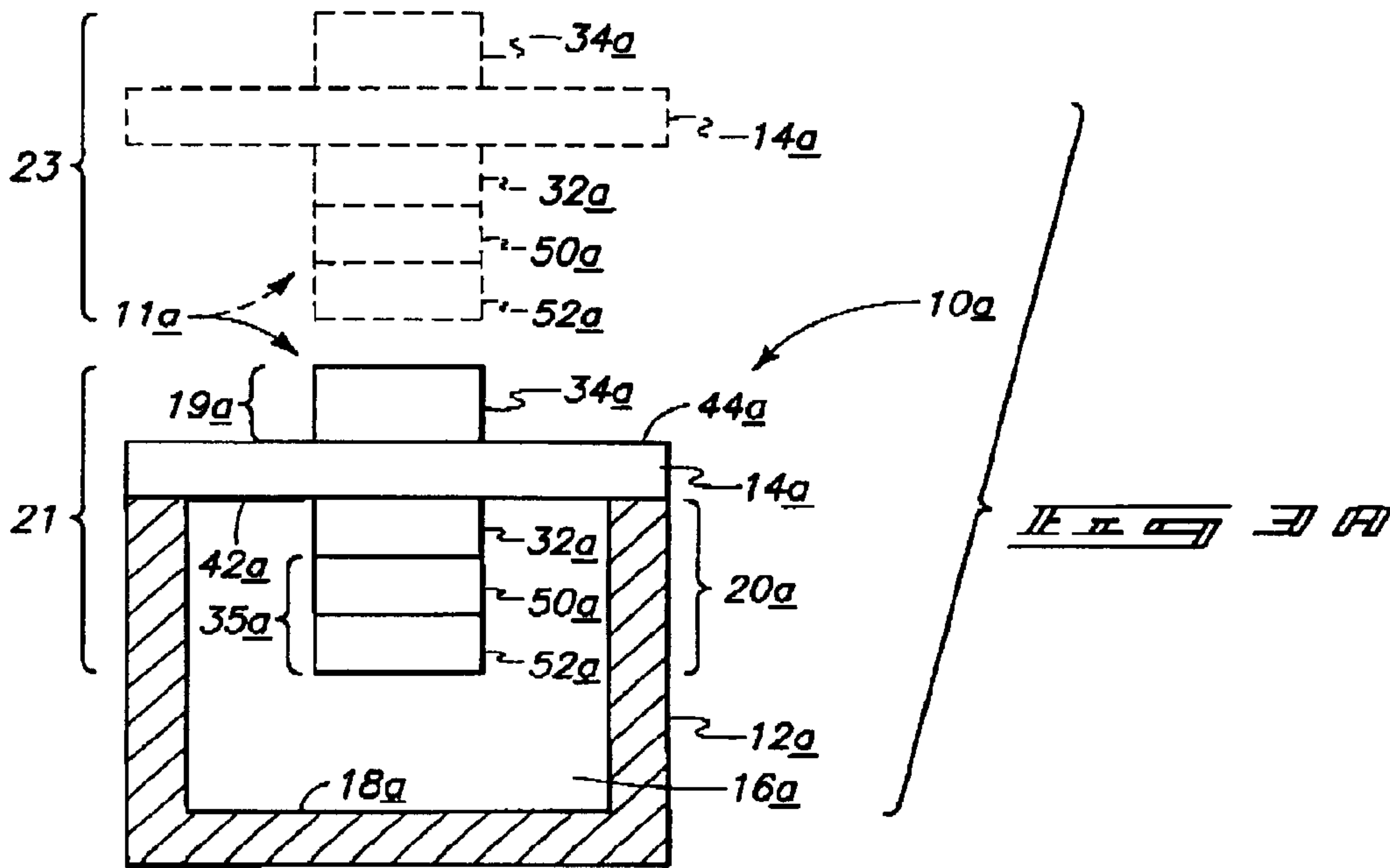
2002/0125425 A1* 9/2002 Kato 250/288
 2006/0219898 A1* 10/2006 McLuckey et al. 250/288

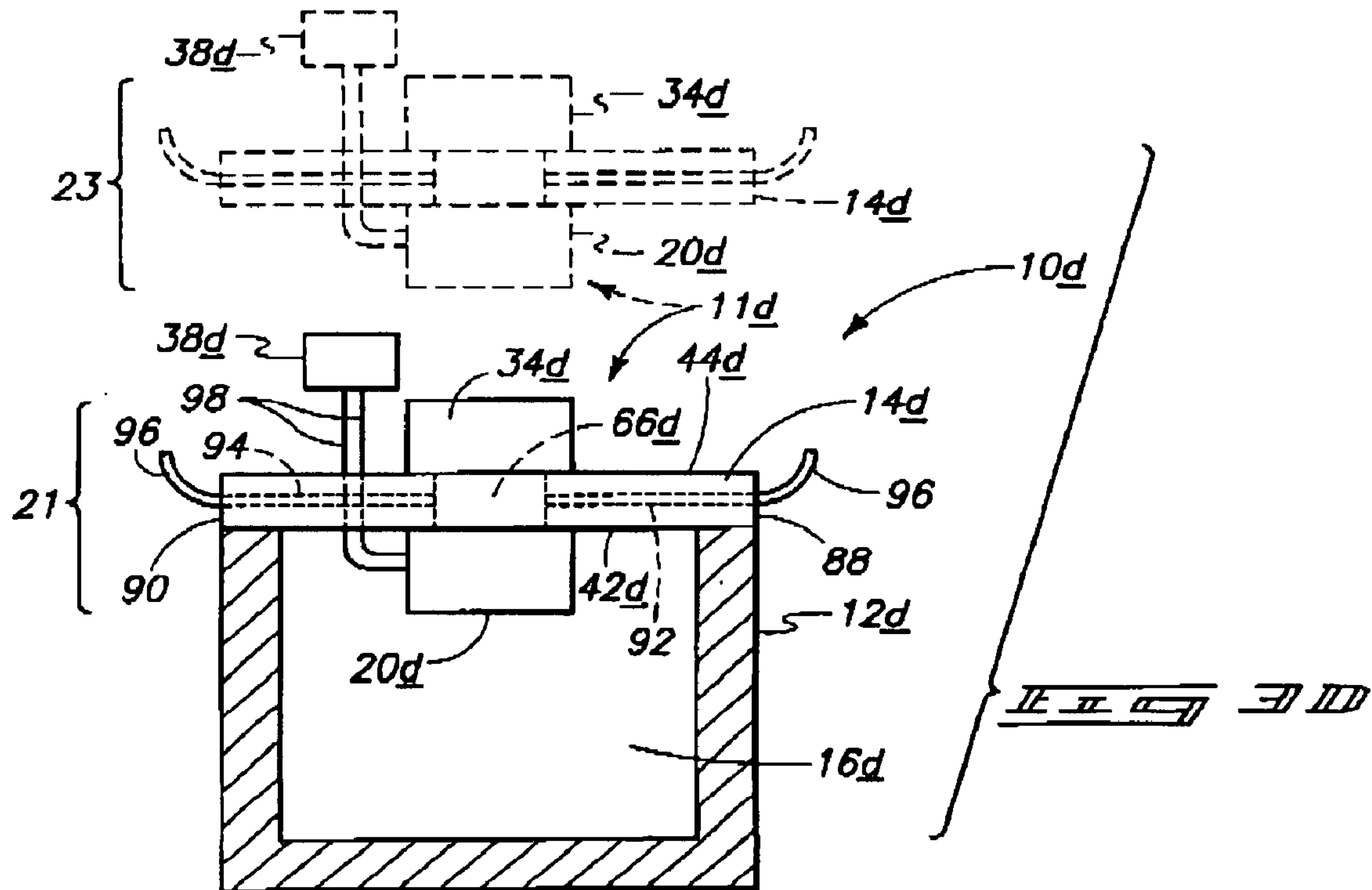
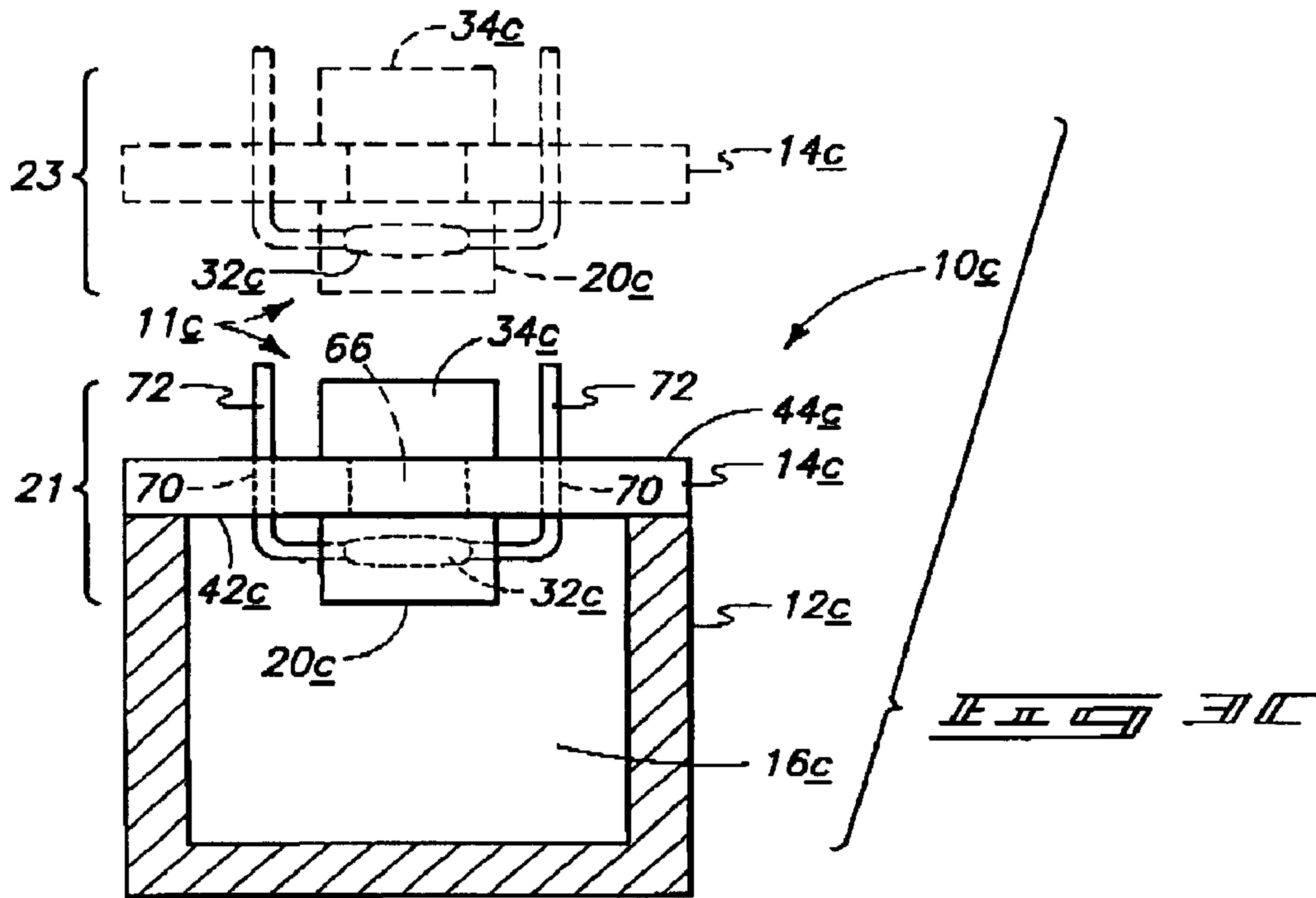
OTHER PUBLICATIONS

Johnson, et al., *Membrane Introduction Mass Spectrometry; Trends And Applications*, Mass Spectrometry Reviews (2000) 19, pp. 1-37.
 Carroll, et al., *A Dual Vacuum Chamber Fourier Transform Mass Spectrometer with Rapidly Interchangeable LSIMS, MALDI, and ESI Sources: Initial Results with LSIMS and MALDI*, Analytical chemistry, vol. 78, No. 10, May 15, 1996, pages.
<http://www.micromass.co.uk>; 3 pps.
 Vestal, Marvin L.; *Methods of Ion Generation*; Chem Rev. 2001, 101(2); pp. 361-375.

* cited by examiner







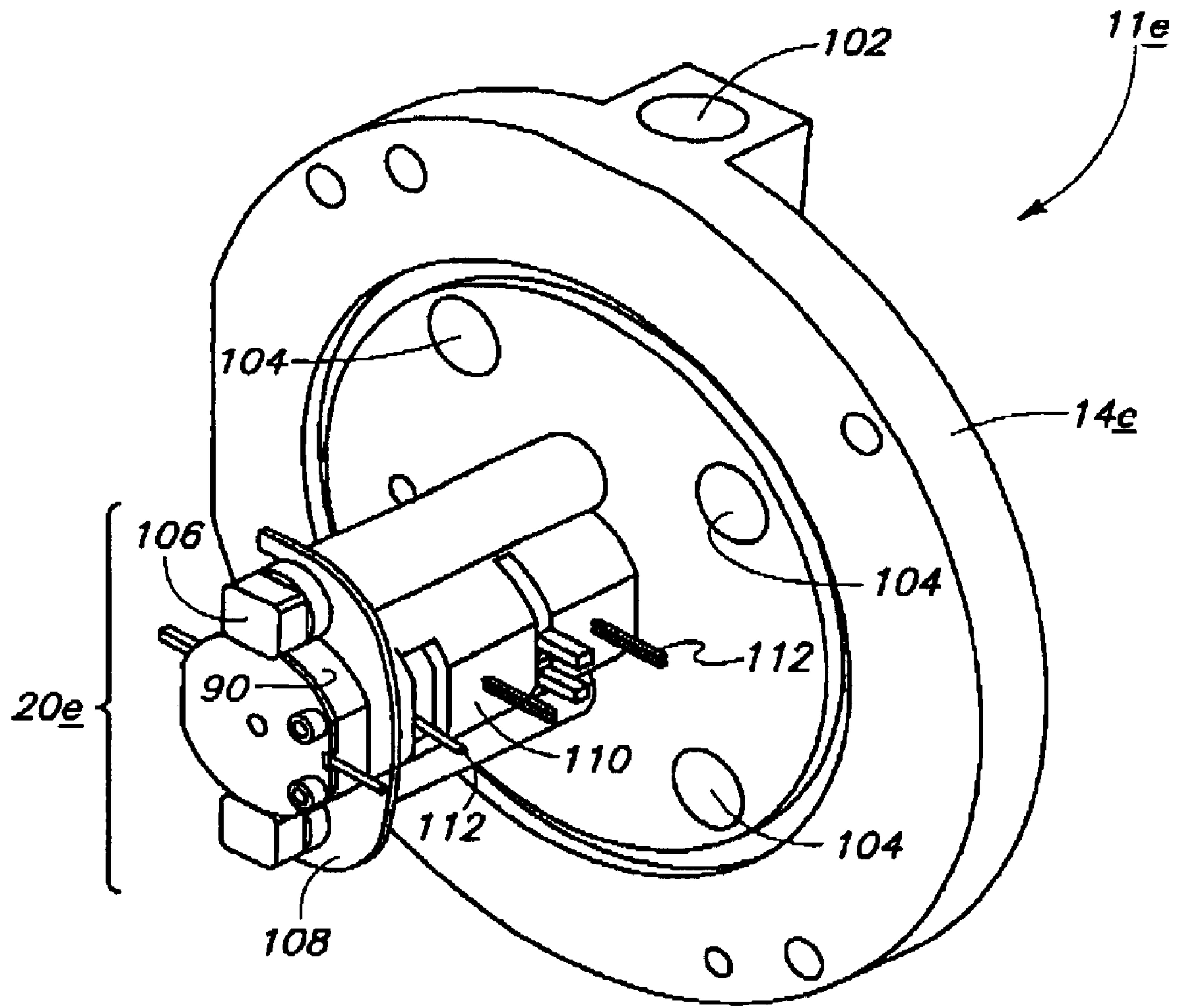


FIG. 11e

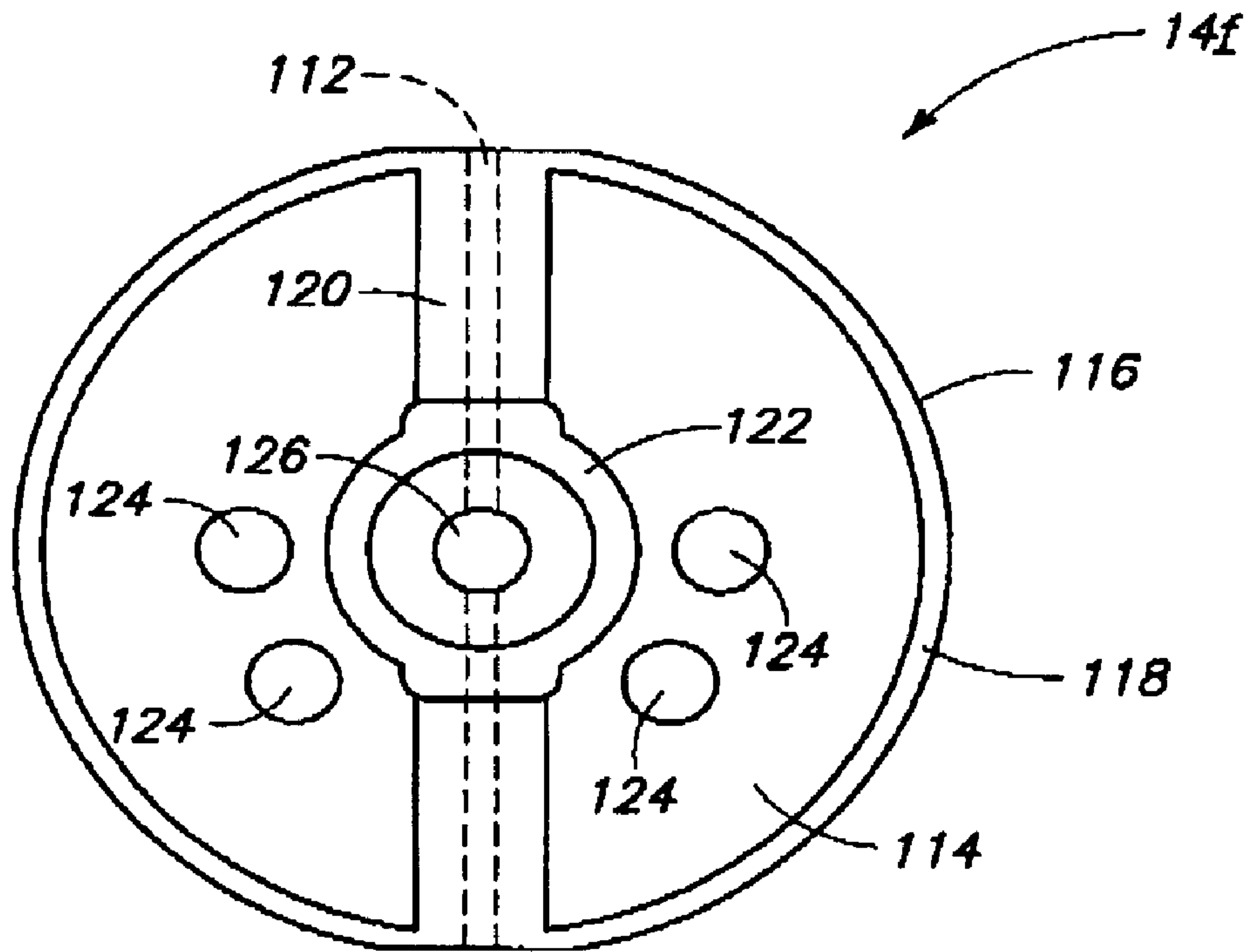


FIG. 5F

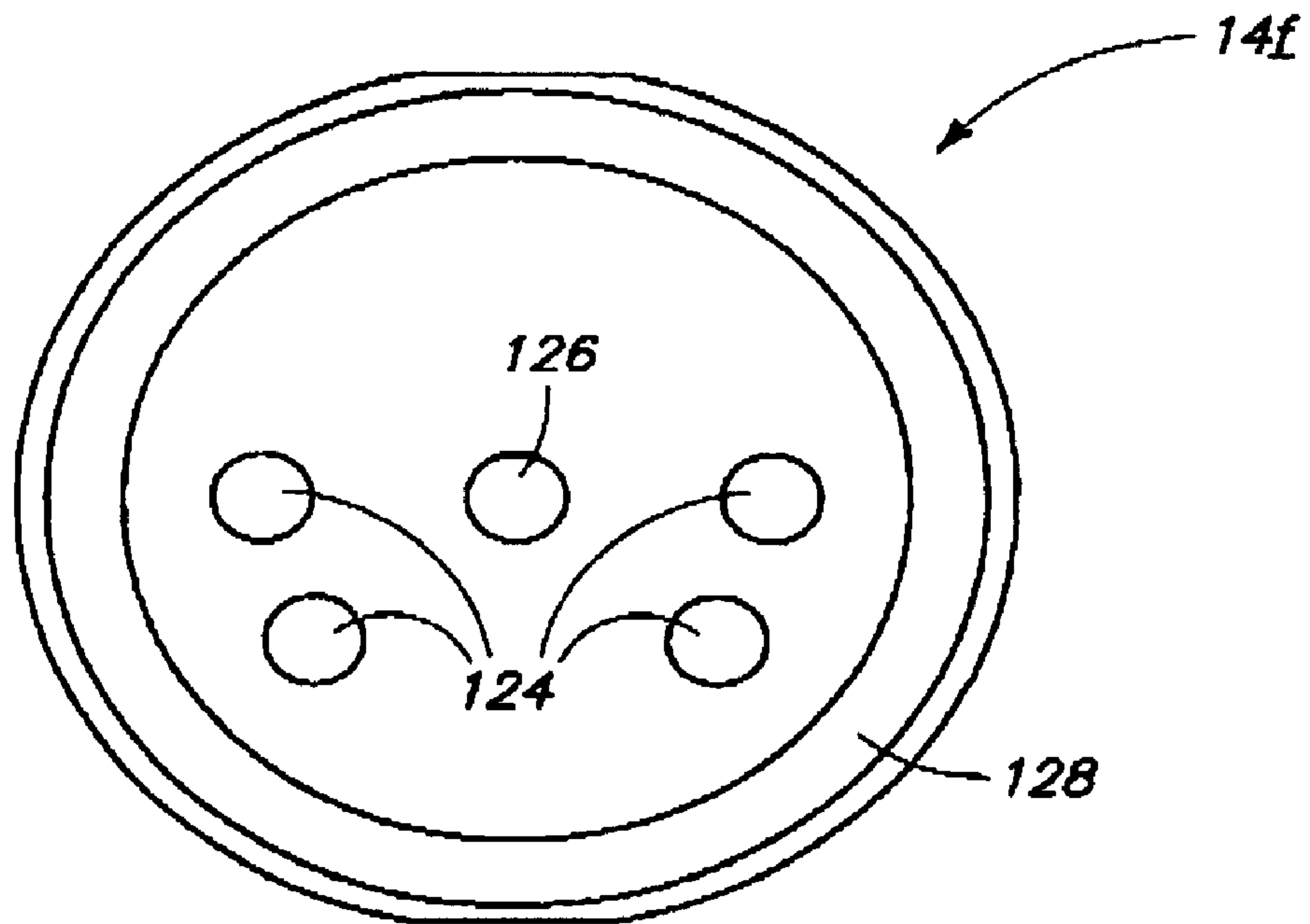
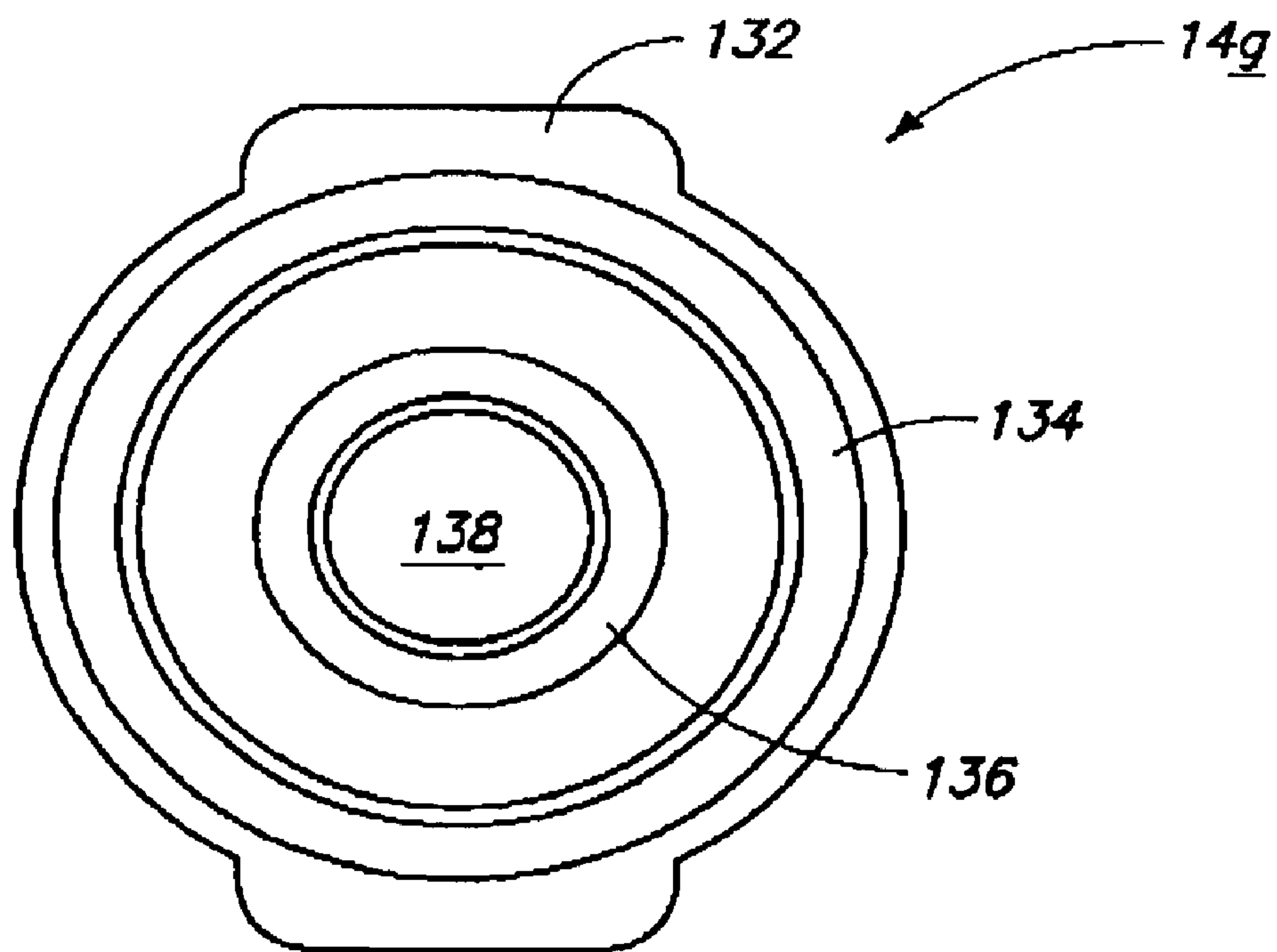
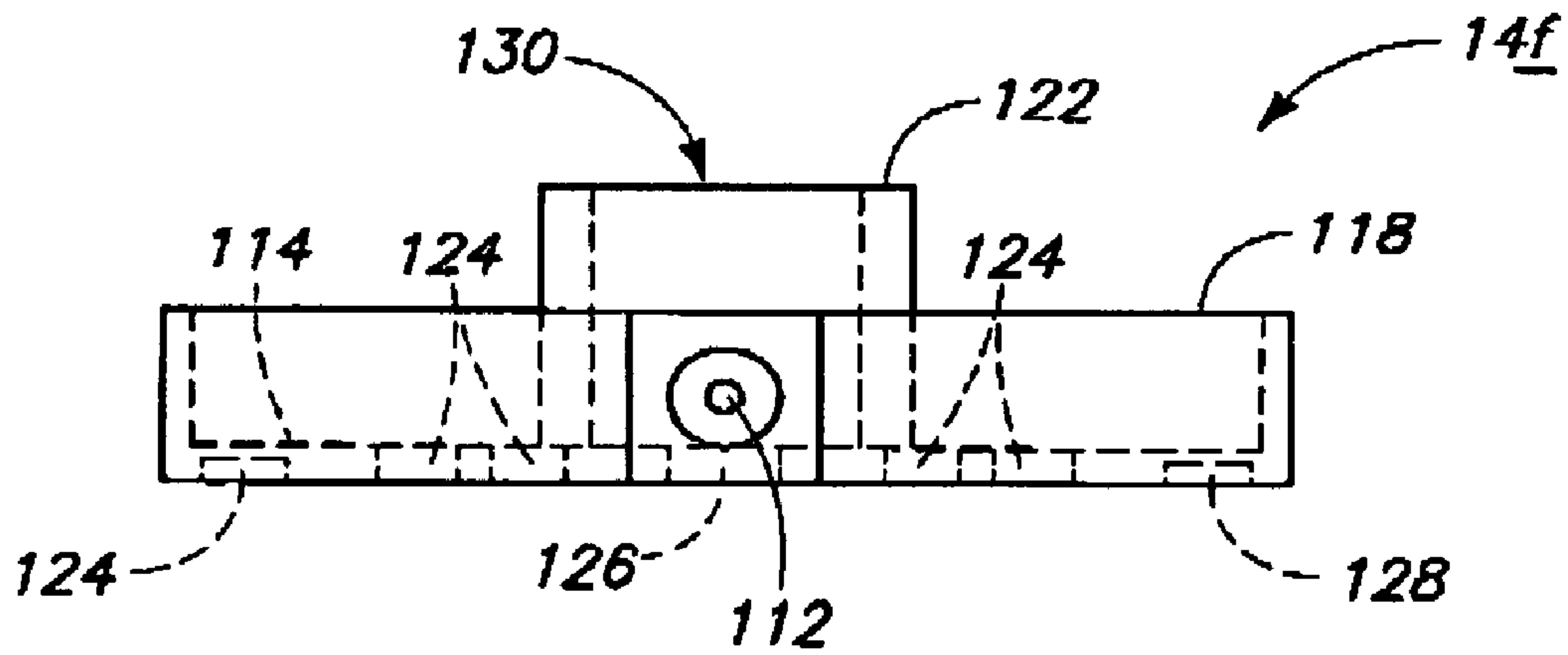


FIG. 5G



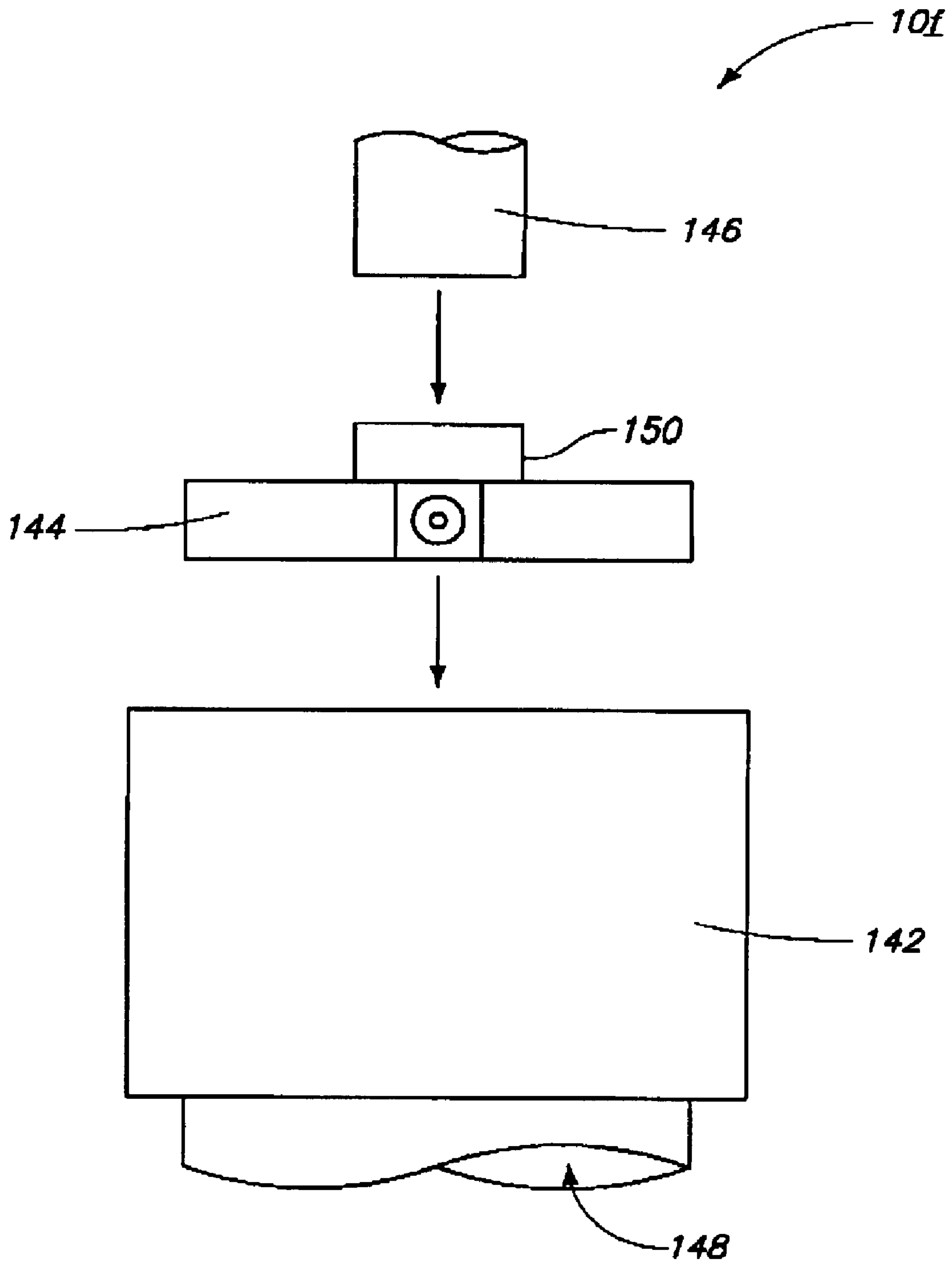
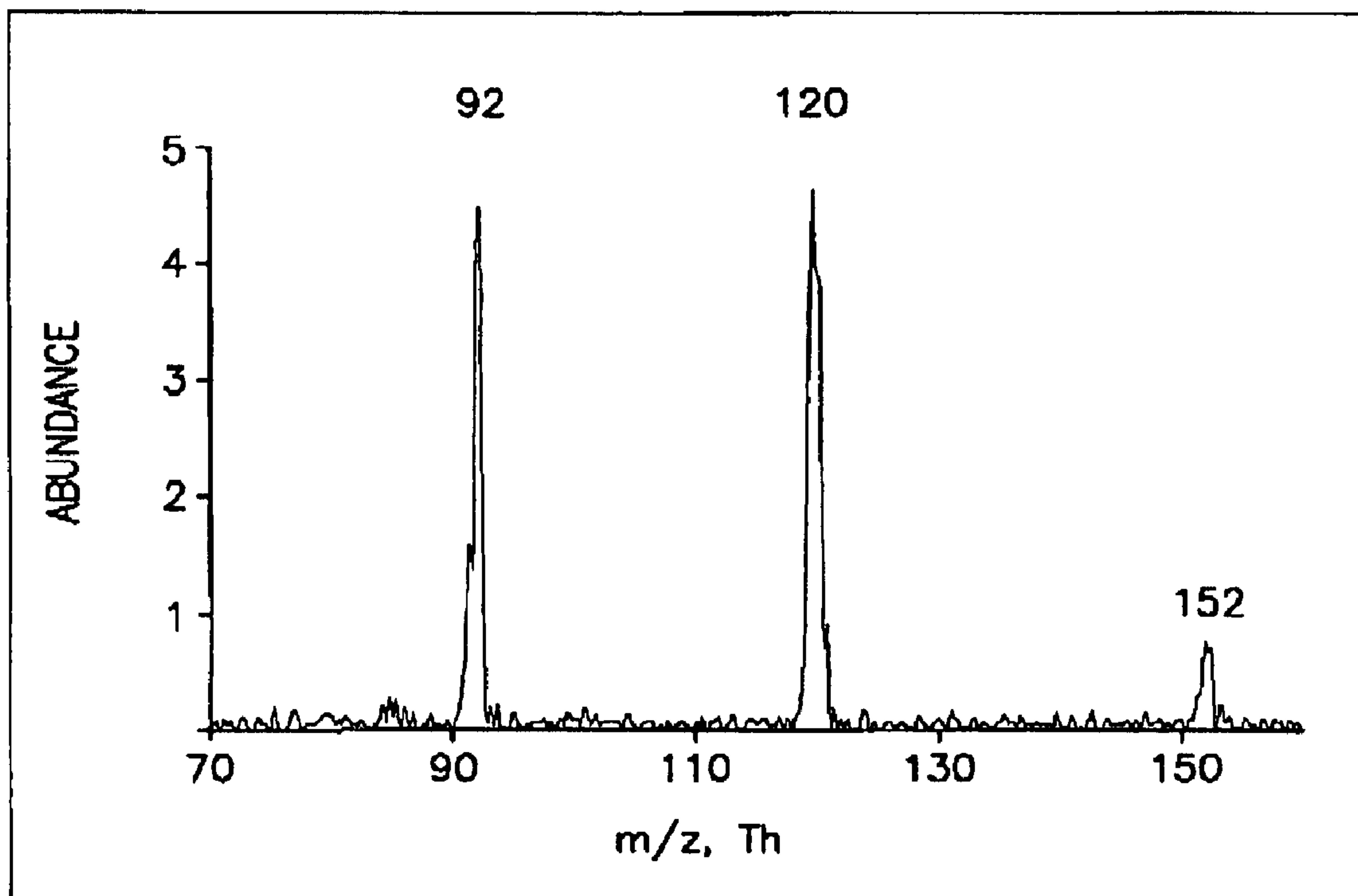


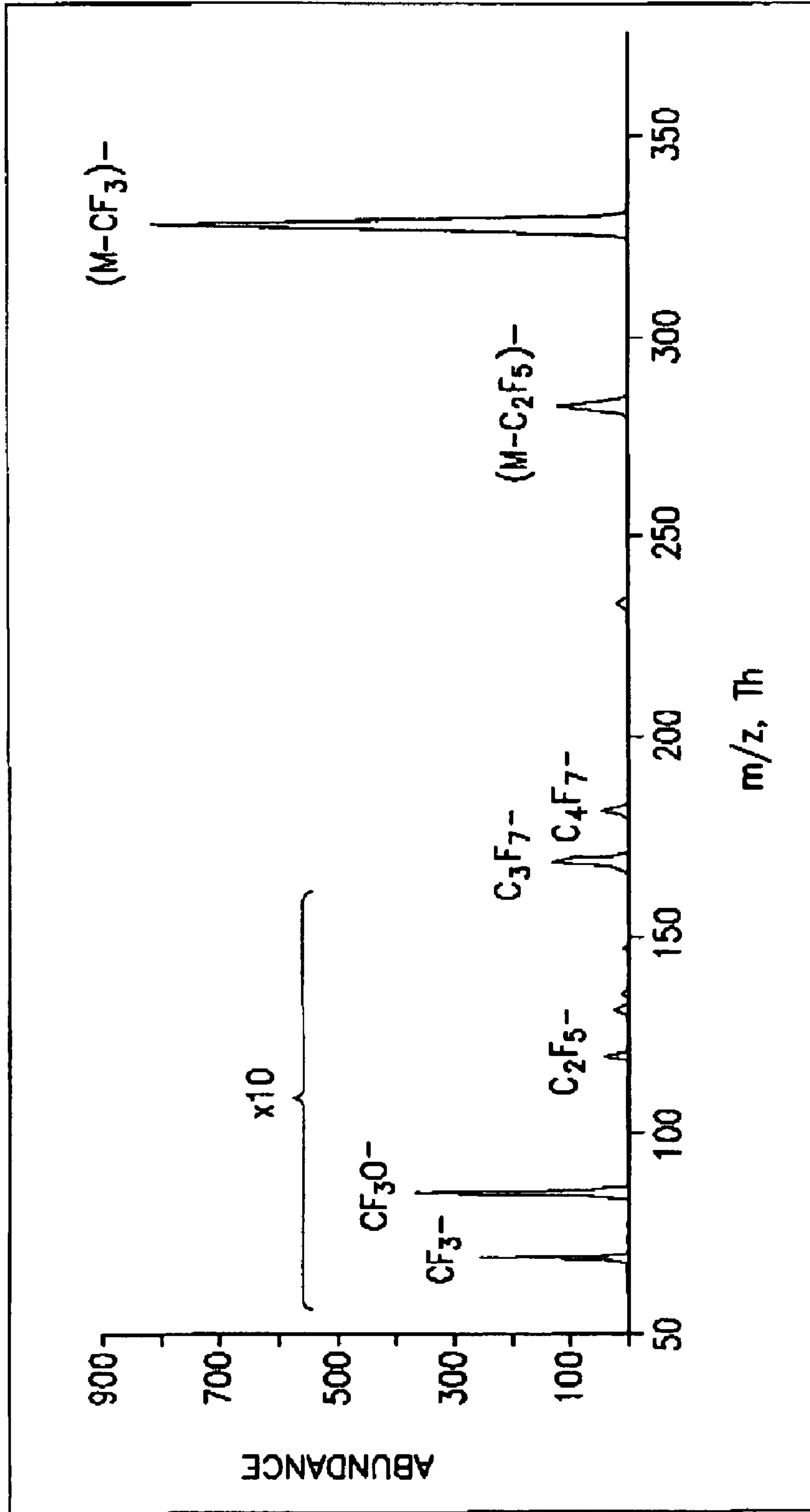
Fig. 10f

SPECTRUM A



JEOL

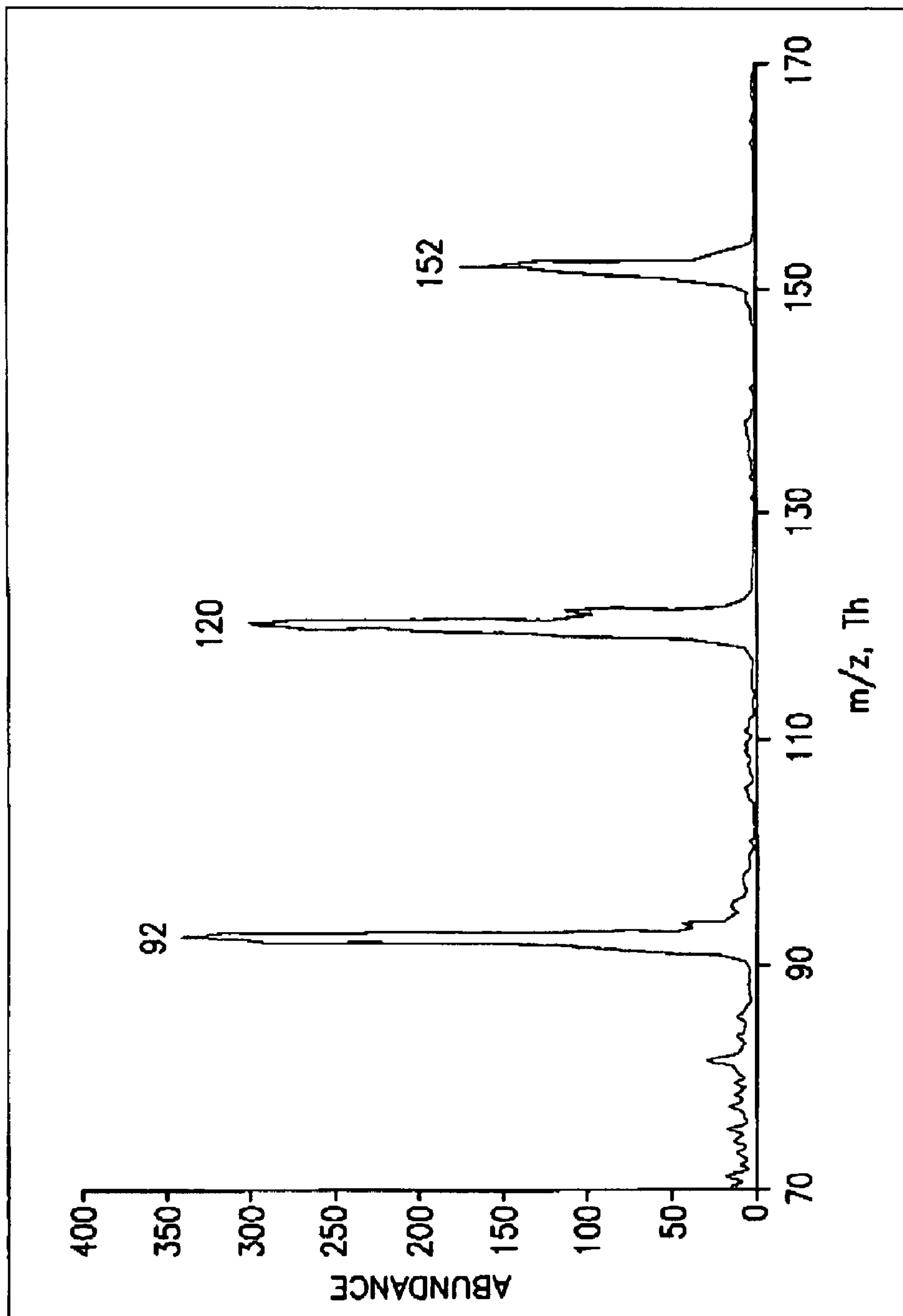
SPECTRUM B



Note: Spectrum was collected in two segments.

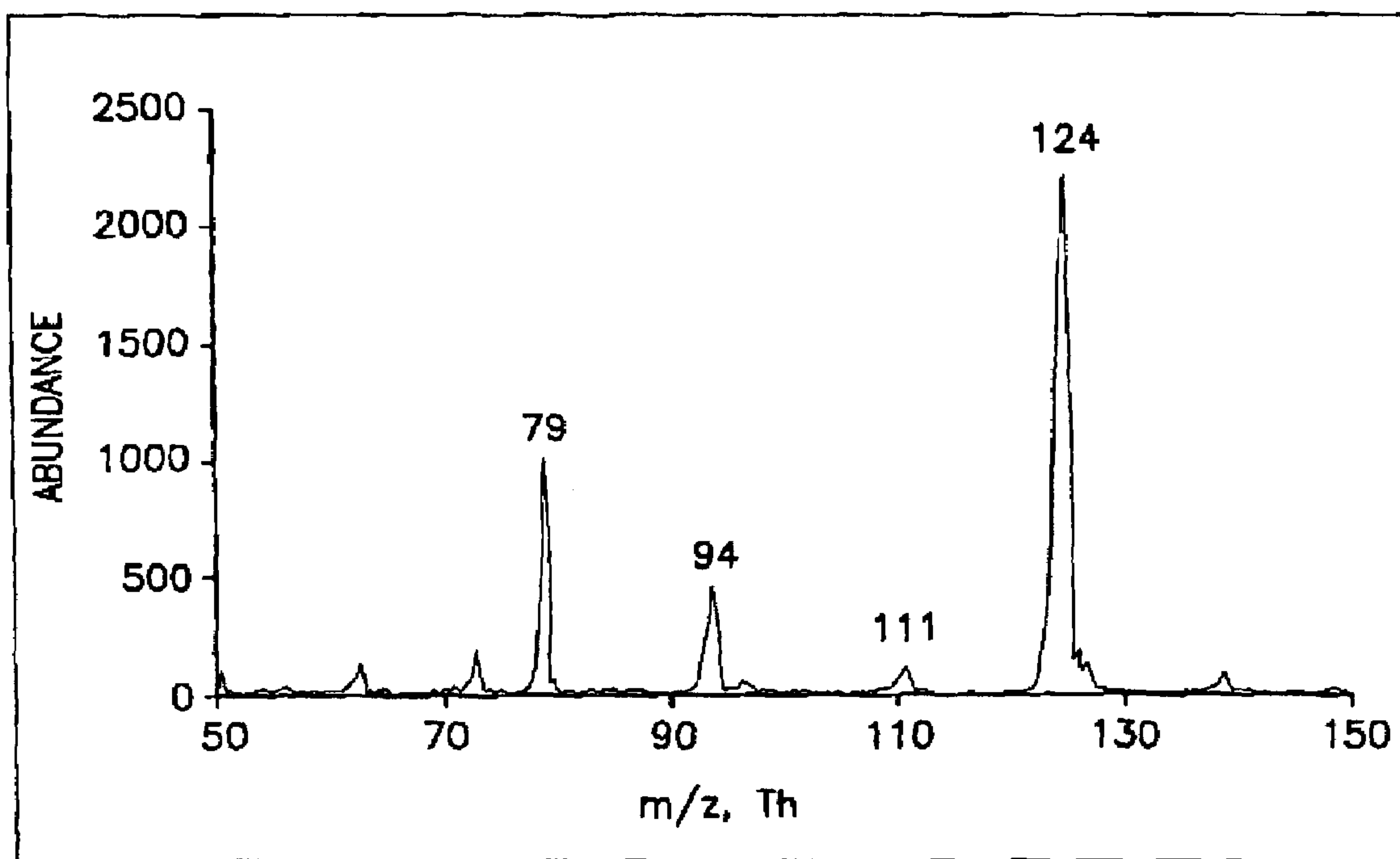
EX 9

SPECTRUM C



EX-100

SPECTRUM D



11 11

1

**MASS SPECTROMETER ASSEMBLIES, MASS
SPECTROMETRY VACUUM CHAMBER LID
ASSEMBLIES, AND MASS SPECTROMETER
OPERATIONAL METHODS**

RELATED PATENT DATA

This application is a 35 U.S.C. §371 of and claims priority to PCT International Application No. PCT/US04/01144, which was filed 16 Jan. 2004 (16.01.04), and was published in English, which claims priority under 35 U.S.C. §119 to U.S. Provisional Patent Application Ser. No. 60/440,887 filed Jan. 17, 2003 entitled "Interchangeable Mass Spectrometer Inlet/Ionization Source", the entirety of both of which are incorporated by reference herein.

TECHNICAL FIELD

The present disclosure relates generally to chemical analysis and more particularly to mass spectrometer assemblies, mass spectrometry vacuum chamber lid assemblies, and mass spectrometer operational methods.

BACKGROUND ART

Characterization of compounds utilizing mass spectrometry and varying sources of ionization is well accepted in the field of analytical chemistry as a technique that allows for the further elucidation of analytes and their specific chemistries. However, mass spectrometer instrumentation is costly and because most labs are unable to configure many instruments with unique ionization sources, analysts are typically required to configure one instrument with a single source and then reconfigure with different sources as analysis dictates. This change-out between sources can be problematic, particularly since mass spectrometer instrumentation must be configured under a vacuum and nanogram quantities of contaminant materials can provide background noise rendering the instrument practically useless.

SUMMARY

Mass spectrometer assemblies are provided that can include in one embodiment: a base configured to define at least a portion of a vacuum chamber volume within which at least some operations may be performed with respect to mass spectrometry; a mass separator component configured to perform at least some operations with respect to mass spectrometry within the vacuum chamber volume; a lid coupled to the mass separator component and configured to be removably operably coupled with respect to the base; and wherein the lid is configured to be positioned in a first operable position to form a hermetical seal with the base and provide the mass separator component with the vacuum chamber volume and a second operable position wherein at least a portion of the lid is spaced from the base and the mass separator component is at least partially removed from the vacuum chamber volume.

Mass spectrometry vacuum chamber lid assemblies are provided that can include, in one embodiment, a body having an interior surface coupled to a mass separator component, wherein the body is configured to at least partially define a volume partially surrounding the mass separator component when the body is hermetically sealed to a housing of a vacuum chamber assembly, wherein the body is further configured to be removable from the vacuum chamber volume to at least partially remove the mass separator component from the vacuum chamber volume.

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Mass spectrometer operational methods are provided that can include, in one embodiment: providing a mass spectrometry assembly comprising a base and a lid, the base and lid substantially defining a vacuum chamber volume when the lid is affixed to the base in a position operable to perform at least some operations with respect to mass spectrometry, wherein a mass separator component is coupled to the lid and occupies a portion of the vacuum chamber volume in the position; first performing mass analysis using the mass spectrometry assembly in the position; after the first performing, at least partially removing the lid from the base, wherein the at least partially removing of the lid also at least partially removes the mass separator component from the vacuum chamber volume; inspecting the mass separator component with the mass separator component removed from the vacuum chamber volume; returning the lid to the position after the inspecting; and second performing mass analysis using the mass separator after the returning.

BRIEF DESCRIPTION OF THE DRAWINGS

Preferred embodiments of the invention are described below with reference to the following accompanying drawings.

FIG. 1 is an illustrative representation of a mass spectrometry assembly according to an embodiment.

FIG. 2 is a block diagram of mass spectrometry components according to an embodiment.

FIG. 3a is an illustrative representation of a mass spectrometry assembly according to an embodiment.

FIG. 3b is an illustrative representation of a mass spectrometry assembly according to an embodiment.

FIG. 3c is an illustrative representation of a mass spectrometry assembly according to an embodiment.

FIG. 3d is an illustrative representation of a mass spectrometry assembly according to an embodiment.

FIG. 4 is an isometric view of a mass spectrometry assembly according to an embodiment.

FIG. 5a is a plan view of a mass spectrometry assembly according to an embodiment.

FIG. 5b is a plan view of the mass spectrometry assembly of FIG. 5a according to an embodiment.

FIG. 5c is a side view of the mass spectrometry assembly FIGS. 5a and 5b according to an embodiment.

FIG. 6 is a plan view of a mass spectrometry assembly according to an embodiment.

FIG. 7 is an illustrative representation of a mass spectrometry assembly according to an embodiment.

FIG. 8 is example data acquired utilizing assemblies and methods of the disclosure.

FIG. 9 is example data acquired utilizing assemblies and methods of the disclosure.

FIG. 10 is example data acquired utilizing assemblies and methods of the disclosure.

FIG. 11 is example data acquired utilizing assemblies and methods of the disclosure.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

At least some embodiments provide mass spectrometry assemblies and mass spectrometer operational methods. Exemplary configurations of these assemblies and methods are described with reference to FIGS. 1-11.

Referring first to FIG. 1, a mass spectrometry assembly 10 is shown that comprises a base 12 and a lid 14. Together lid 14 and base 12 can define a volume 16. In an exemplary embodi-

ment lid 14 and base 12 define a vacuum chamber housing of mass spectrometry assembly 10. The vacuum chamber housing can define volume 16 which can be referred to as a vacuum chamber volume in embodiments wherein a vacuum is provided. A vacuum source (not shown) can be in fluid connection to base 12 and provide vacuum to volume 16.

Base 12 can be constructed of a single structure or can be constructed of multiple components. Exemplary components include walls 17 and a bottom 18. In the exemplary configuration of FIG. 1, walls 17 may be a continuous wall of a cylinder. Other geometries or arrangements of base 12 are possible. The structure and or components of base 12 and/or lid 14 can be fabricated of materials such as aluminum, stainless steel, and/or other materials. In exemplary embodiments lid 14 can be referred to as a body or as a flange and/or an adapter. According to exemplary aspects, base 12 can be configured to define at least a portion of volume 16. In exemplary embodiments walls 17 of base 12 can be affixed to lid 14 by a hermetical seal (not shown). An exemplary hermetical seal can include an O-ring arrangement between lid 14 and base 12. An appropriate removable fastener (not shown) may also be provided to maintain lid 14 and base 12 in a sealed arrangement.

In one embodiment, mass spectrometry assembly 10 comprises one or more components configured to perform operations with respect to mass spectrometry analysis, and accordingly, such components may be referred to as mass spectrometry components 30. In one possible implementation, lid 14 is coupled with one or more of components 30. Further, an individual one of components 30 may be internally or externally coupled with lid 14. For example, in the embodiment shown in FIG. 1, lid assembly 11 comprises lid 14 coupled with an external component 19 and an internal component 20. In the illustrated exemplary embodiment, external component 19 is external of a volume 16 and internal component 20 is at least partially within volume 16. An individual one of components 30 may also be provided coupled with lid 14 in other arrangements, for example, internal of or defined by a volume of lid 14.

Within volume 16 at least some mass spectrometry operations can be performed using internal component 20. Some mass spectrometry operations can also be performed using external component 18. In an exemplary aspect, lid 14 and/or lid assembly 11 can be configured to be removably operably coupled with respect to base 12. Lid 14 can be configured to be positioned in a first operable position 21. In position 21, lid 14 can form a hermetical seal with base 12 and provide component 20 within volume 16. In position 21, mass spectrometry assembly 10 can be used to perform at least some operations with respect to analysis of a sample.

Lid 14 can also be configured to be positioned in a second operable position 23. In position 23, at least a portion of lid 14 can be spaced from base 12 and component 20 can be at least partially removed from volume 16. In an exemplary aspect, an entirety of lid 14 can be spaced from base 12 and component 20 can be at least partially removed from volume 16. In another exemplary aspect, at least a portion of lid 14 can be spaced from base 12 and component 20 can be entirely removed from volume 16. In another exemplary aspect, an entirety of lid 14 can be spaced from base 12 and an entirety of component 20 can be removed from volume 16. Second operable position 23 facilitates access to internal equipment 20 in one embodiment. Other operable positions intermediate operable positions 21 and 23 are possible. The plural operable positions of lid 14 and/or lid assembly 11 may refer to an exemplary embodiment of assembly 10 where lid 14 and/or lid assembly 11 are detached and reattached numerous times

with respect to base 12 when used during mass spectrometry operations (e.g. service, reconfiguration, maintenance, etc.).

Referring next to FIG. 2, components 30 of a mass spectrometer according to one embodiment are shown. As represented in FIG. 2, components 30 can include a sample inlet component 32 operationally connected to an ion source component 34 which can be operationally connected to a mass separator component 35 which can be operationally connected to a detector component 36. These general components can be operationally connected to a processing and control device component 38. Exemplary embodiments provide for the use of components 30 to perform mass spectrometry. Components 30 can be operationally connected as shown in FIG. 2 or operationally connected in other configurations enabling mass spectrometry operations. Further, other arrangements including more or less or alternative components are possible.

As depicted in FIG. 2, a sample 40 can be introduced into sample inlet component 32. For purposes of this disclosure, sample 40 represents any chemical composition including both inorganic and organic substances in solid, liquid and/or vapor form. Specific examples of sample 40 suitable for analysis include volatile compounds such as toluene or other specific examples including highly-complex non-volatile protein based structures such as bradykinin. In certain aspects, sample 40 can be a mixture containing more than one substance or in other aspects sample 40 can be a substantially pure substance. Analysis of sample 40 can be performed according to exemplary aspects described below.

Sample inlet component 32 can be configured to introduce an amount of sample 40 into assembly 10 (FIG. 1) for analysis. Depending upon sample 40, sample inlet component 32 may be configured to prepare sample 40 for ionization. Types of sample inlet components 32 can include batch inlets, direct probe inlets, chromatographic inlets, and permeable, semi-permeable, solid phase microextraction (SPME), and/or capillary membrane inlets. Sample inlet component 32 can also include means for preparing sample 40 for analysis in the gas, liquid and/or solid phase. In some aspects, sample inlet component 32 may be combined with ion source component 34.

Ion source component 34 can be configured in exemplary embodiments to receive sample 40 directly or in other exemplary embodiments to receive sample 40 from sample inlet component 32. Ion source component 34 can be configured to convert portions or an entirety of sample 40 into analyte ions in one example. This conversion can include the bombardment of sample 40 with electrons, ions, molecules, and/or photons. This conversion can also be performed by thermal or electrical energy.

Ion source component 34 may utilize, for example, electron ionization (EI, typically suitable for the gas phase ionization), photo ionization (PI), chemical ionization, collisionally activated dissociation and/or electrospray ionization (ESI). For example in PI, the photo energy can be varied to vary the internal energy of the sample. Also, when utilizing ESI, sample 40 can be energized under atmospheric pressure and potentials applied when transporting ions into volume 16 of exemplary mass spectrometer assembly 10 (FIG. 1) can be varied to cause varying degrees of dissociation.

The analyte ions can proceed to mass separator component 35. Mass separator component 35 can include one or more of linear quadrupoles, triple quadrupoles, quadrupole ion traps (Paul), cylindrical ion traps, linear ion traps, rectilinear ion traps, ion cyclotron resonance, quadrupole ion trap/time-of-flight mass spectrometers, or other structures. Mass separator component 35 can also include focusing lenses as well as tandem mass separator components such as tandem ion traps

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or ion traps and quadrupoles in tandem. In one implementation at least one of multiple tandem mass separator components can be an ion trap. Tandem mass separator components can be placed in series or parallel. In an exemplary implementation, tandem mass separator components can receive ions from the same ion source component. In an exemplary aspect the tandem mass separator components may have the same or different geometric parameters. The tandem mass separator components may also receive analyte ions from the same or multiple ion source components.

Analytes may proceed to detector component 36. Exemplary detector components include electron multipliers, Faraday cup collectors, photographic and scintillation-type detectors. The progression of mass spectrometry analysis from sample inlet component 32 to detector component 36 can be controlled and monitored by a processing and control device component 38.

Acquisition and generation of data can be facilitated with processing and control device component 38. Processing and control device component 38 can be a computer or mini-computer or other appropriate circuitry that is capable of controlling components 30. This control can include for example the specific application of voltages to ion source component 34 and mass separator component 35, as well as the introduction of sample 40 via sample inlet component 32 and may further include determining, storing and ultimately displaying mass spectra recorded from detector component 36. Processing and control device component 38 can contain data acquisition and searching software. In one aspect such data acquisition and searching software can be configured to perform data acquisition and searching that includes the programmed acquisition of total analyte count. In another aspect, data acquisition and searching parameters can include methods for correlating the amount of analytes generated to pre-determine programs for acquiring data.

Referring again to FIG. 1, individual ones of the general components of a mass spectrometer may be positioned as an internal component 20 or as an external component 19 as desired by those of ordinary skill in the art. For example, in different applications or configurations, individual ones of general components 32-26 may be arranged as internal or external components as desired. In one embodiment, internal component(s) 20 can include one or more of sample inlet component 32, ion source component 34, mass separator component 35 and/or the detector component 36 in various configurations to perform mass spectrometry. In one embodiment, external component(s) 19 can include one or more of sample inlet component 32, ion source component 34 and/or processing and control device 36.

In an exemplary embodiment, internal and/or external components include multiple components such as multiple ion source components. These multiple components can be configured as external, internal or external and internal components.

Exemplary arrangements of the mass spectrometry components and lid assemblies are shown in FIGS. 3a-3d.

Referring to FIG. 3a, an exemplary mass spectrometer assembly 10a is shown that includes lid 14a, base 12a and volume 16a. Lid 14a can include an interior surface 42a and an exterior surface 44a. Exemplary embodiments include lid assembly 11a that can include external component 19a including ion source component 34a coupled to external surface 44a. In the exemplary depiction, lid assembly 11a can include internal components 20a including sample inlet component 32a and mass separator component 35a including focusing lenses 50a and an ion trap 52a coupled to internal surface 42a to process analytes in a direction that is substan-

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tially perpendicular to the alignment of inner surface 42a toward the detector component (not shown). For example, analytes can be processed in a direction towards bottom 18a of base 12a. Sample inlet component 32a can be located between mass separator component 35a and internal surface 42a. Sample inlet component 34a and mass separator component 35a can be in a substantially stacked configuration below and coupled to inner surface 42a.

Referring to FIG. 3b, an exemplary mass spectrometry assembly 10b is shown that includes ion source component 34b coupled to external surface 44b. Lid assembly 11b can include lid 14b coupled to one or both of external component 19b and/or internal component 20b. As exemplarily depicted, internal components 20b include sample inlet component 34b and mass separator components 35b including focusing lenses 50b and ion trap 52b coupled to internal surface 42b to process analytes in a direction that is substantially parallel to the alignment of inner surface 42b toward the detector component (not shown). For example, analytes can be processed in a direction towards wall 17b of base 12b.

Referring again to FIGS. 3a and 3b, in operable position 21, internal components 20a-b including sample inlet component 34a-b, focusing lenses 50a-b and ion trap 52a-b and detector (not shown) can be within volume 16a-b. In an exemplary embodiment volume 16a-b can at least partially surround internal components 20a-b when lid 14a-b is in operable position 21. In other exemplary aspects internal components 20a-b can also include ion source component 34a-b.

In operable position 23, one or more of internal components 21a-b including sample inlet component 34a-b, focusing lenses 50a-b, and ion trap 52a-b can be at least partially removed from volume 16a-b. In an exemplary aspect, in operable position 23, one or more of the internal components can be entirely removed from volume 16a-b. For example and by way of example only, in operable position 23: sample inlet component 32a-b can be entirely removed from volume 16a-b while mass separator component 35a-b is not removed; sample inlet component 34a-b and focusing lenses 50a-b can be entirely removed from volume 16a-b while ion trap 52a-b is not removed; sample inlet component 32a-b and ion trap 52a-b can be entirely removed while focusing lenses 50a-b are not removed; both sample inlet component 32a-b and mass separator component 35a-b including both focusing lenses 50a-b and ion trap 52a-b can be entirely removed; focusing lenses 50a-b can be entirely removed while ion trap 52a-b is not removed; and/or ion trap 52a-b can be entirely removed while focusing lenses 50a-b are not removed from volume 16a-b.

Referring next to FIG. 3c, another exemplary mass spectrometry assembly 10c is shown and includes base 12c and lid 14c. As shown in FIG. 3c, lid 14c can include an interior surface 42c, an exterior surface 44c, and an opening 66 extending from interior surface 42c to exterior surface 44c. In the exemplary Figure, ion source component 34c is coupled to exterior surface 44c. In an exemplary aspect ion source component 34c can be coupled to exterior surface 44c in fluid communication with opening 66. In the exemplary depiction, internal components 20c can be coupled to interior surface 42c. In an exemplary aspect internal components 20c can couple to interior surface 42c in fluid communication with opening 66. Exemplary aspects include providing fluid communication between external components 19c and internal components 20c via opening 66. In an exemplary embodiment, ion source component 34c is in fluid communication with sample inlet component 32c through opening 66. Lid

assembly **11c** can include lid **14c** having opening **66** and one or both of internal components **20c** and external components **19c**.

Lid **14c** can also be configured to provide sample **40** (FIG. 2) to sample inlet component **32c**. In an exemplary embodiment openings **70** can be provided that extend into volume **16c** from outside volume **16c** through lid **14c**. Sample tubing **72** can be provided through or as part of openings **70** and configured to provide sample **40** to sample inlet component **32c**. In first operable position **21** internal component(s) **20c** including sample inlet component **32c** can be within volume **16c**. In second operable position **23**, portions or an entirety of one or more internal component(s) **20c** which can include sample inlet component **32c** can be removed from volume **16c**.

Referring next to FIG. 3d, mass spectrometry assembly **10d** is shown including lid **14d** having an interior surface **42d** and an exterior surface **44d** having opening **66d** therethrough, and edges **88** and **90** extending between interior surface **42d** and exterior surface **44d**. In an exemplary embodiment, lid **14d** includes openings **92** and **94** extending from edge **88** and edge **90** respectively to and in fluid communication with opening **66d**. In an exemplary aspect openings **92** and **94** can provide fluid communication between outside of volume **16d** and opening **66d**. Conduit or tubing **96** can be configured to facilitate fluid communication to opening **66d** through openings **92** and **94**. Embodiments of lid **14d** also include single openings extending from a single edge such as edge **88** and opening **92** to opening **66d**. Embodiments of lid **14d** also include openings (not shown) extending from outside volume **16d** such as exterior surface **88** to opening **66**. Openings **92** and **94** can be utilized in exemplary aspects to: provide auxiliary pumping to opening **66d**; to provide reagent gas such as chemical ionization reagent gases to opening **66d** during ionization using ion source **34d**; and/or to provide sample **40** to opening **66d**. In an exemplary embodiment, opening **92** can serve as an inlet port while opening **94** may serve as an outlet port for samples circulating through tubing within lid **14d**. Openings **92** and/or **94** can also provide for gas pumping on central opening **66d** should an intermediate pressure be needed for proper operation of ion source component **34d** or in the event of a large sample influx such as is the case with liquid samples.

Lid **14d** can also be configured to provide control and/or power to internal component **20d** for example through electrical wiring **98**. Electrical wiring **98** can be incorporated as part of lid **14d** or through openings provided in lid **14d**. Electrical wiring **98** can be configured to control internal component(s) **20d** such as sample inlet component **32** and mass separator component **35** from processing and control device component **38**. In first operable position **21** internal component(s) **20d** and at least some wiring **98** can be within volume **16d**. In second operable position **23**, portions or an entirety of one or more internal component(s) **20d** and wiring **98** can be removed from volume **16d**. Exemplary embodiments provide for lid assembly **11d** that includes lid **14d** and one or both of internal component **19d** and/or internal component **20d**.

Referring next to FIG. 4, an exemplary mass spectrometer vacuum chamber lid assembly **11e** is shown in isomeric view that includes a lid **14e** and internal components **20e** which may comprise mass separator components. Assembly **11e** may be used in the arrangements of FIGS. 1 and 3 in exemplary configurations. Components **20e** can be coupled to lid **14e** via one or more of a mounting rod **106** and a retainer clip **108**. Mass separator components **20e** can be confined within mass separator housings **110** and can be controlled via electrical signals provided to electrical connections **112**. In one embodiment, mass separator housing **110** includes insulating material. As shown in FIG. 4, lid **11e** includes openings **104**

(see e.g., opening **70**, FIG. 3c) and **102** (see e.g., opening **92**, FIG. 3d). In an exemplary aspect openings **104** can be used to pass wiring, sample, and/or other mass spectrometry components into volume **16** (FIG. 1). Opening **102** may be used as described above with reference to opening **92**.

Referring next to FIGS. 5a-5c an exemplary embodiment of lid **14f** is shown in differing views. Referring first to FIG. 5a an exterior view of lid **14f** is shown having recessed portions **114** radially inward of edges **116** providing an elevated lip **118**. Recessed portion **114** can also be referred to as a groove. Openings **124** are provided through lid **14f** in an exemplary embodiment to provide for feed through connections of electrical wiring and/or sample inlet components. Lid **14f** can also include elevated portions **120**, in an exemplary embodiment bisecting lid **14f**. Opening **126** can be provided through the center of lid **14f** to facilitate, in an exemplary aspect, the introduction of ions, samples or other materials provided during mass spectrometry. Lid **14f** can also include adaptive portions **122** to facilitate the coupling of external components **19** (not shown) such as ion source component **34**. In an exemplary embodiment opening **112** can extend bisecting lid **14f** and through opening **126** to provide fluid communication between edge **116** and opening **126**.

Referring next to FIG. 5b, internal surface view of body **14f** is shown that includes a recessed portion **128** that can facilitate the hermetical sealing of lid **14f** to base **12** of a mass spectrometry assembly **10** (FIG. 1). Referring next to FIG. 5c, a side view of lid **14f** is shown. As shown lip **118** and adaptive portions **122** can extend above recessed surface **114**.

Referring next to FIG. 6, an external view of an exemplary embodiment of lid **14g** is shown that includes recessed groove **134** as well as a mating surface **132** to facilitate sealing lid **14g** to base **12** (FIG. 1). In an exemplary embodiment, base **12** can include a complimentary mating surface (not shown) to mating surface **132**. Exemplary embodiments of lid **14g** can also include a circular plate **136** that in certain aspects can be installed within opening **138** to maintain a pressure in opening **126** (FIGS. 5a-c) that is higher than that within volume **16** (FIG. 1).

Referring to FIG. 7, a mass spectrometry assembly **10f** is shown that includes a base **142**, a lid **144** and an external component **146**. As shown, base portion **142** also includes an opening **148** that in an exemplary embodiment allows access to vacuum pumping and/or detection of ions separated utilizing assembly **10f**. As exemplarily depicted in FIG. 7, external component **146** can include ion source component **34** and in one embodiment can be removably operably coupled with respect to lid **144** and configured to be positioned in a first operable position (not shown) to seal with lid **144** and a second operable position where at least a portion of external component **146** is spaced from lid **144**.

Other aspects provide for the configuration of assembly **10f** with multiple components. Multiple ion sources can be configured to couple with lid **144** in one embodiment. In an exemplary aspect, different ion sources can be configured to be exchanged and/or replaced with respect to assembly **10f**. In an exemplary embodiment, an electron impact ion source may be replaced with a chemical ionization ion source.

Referring to the figures discussed above, mass spectrometer operational methods are also provided that include first performing mass analysis using mass spectrometry assembly **10** in operable position **21**. This performance can include providing sample **40** to volume **16** as described above.

According to an exemplary aspect, mass analysis can include providing ions to the vacuum chamber volume through opening **66** (FIG. **3c**) and contacting the ions with sample **40**. Contacting sample **40** with the ions can occur within opening **66**. In an exemplary embodiment mass analysis can include providing a chemical ionization plasma and a chemical ionization reagent gas, for example through openings **92**, to opening **66d** (FIG. **3d**) to produce ions and exposing sample **40** to the ions to form analytes.

After performing mass analysis, lid **14** can be moved to second operable position **23**. In an exemplary aspect lid **14** can be at least partially removed from base **12** and internal component **20** can be at least partially removed from volume **16**. During mass analysis, components **30** (FIG. **2**) may require inspection between analysis for example for maintenance and trouble shooting requirements.

Internal components **20** such as mass separator components **35** (FIG. **2**) can be inspected with lid **14** in second operable position **23**. In an exemplary embodiment ion source component **34**, such as internal ion source components, including, in exemplary aspects, electron impact filaments, can become fouled during mass analysis and with lid **14** in second operable position **23**, ion source components such as the filaments can be replaced with clean filaments or replacement filaments. In an exemplary embodiment mass separator component **35** such as ion trap **52a-b** (FIGS. **3a-b**) can become fouled or require replacement during mass analysis and with lid **14** in second operable position **23** mass separator component **35** such as ion trap **52a-b** can be replaced with a clean or conditioned ion trap. In an exemplary aspect sample inlet component **34**, such as a semi-permeable membrane can require cleaning and or replacement during mass analysis and with lid **14** in second operable position **23** sample inlet component **32** can be cleaned or replaced.

In an exemplary embodiment, before moving the lid to operable position **23**, ion source component **146** (FIG. **7**) can be at least partially or entirely removed from lid **144** and inspected. Upon inspection, ion source component **146** can be cleaned, replaced, or otherwise manipulated.

After inspection, the lid can be returned to first operable position (not shown) and mass analysis can be performed using components **30** (FIG. **2**).

At least one arrangement facilitates servicing and reconfiguration of assembly **10**. For example, upon removal of the lid assembly or the lid from the base, the internal components, wiring, and tubing, may be removed from the vacuum chamber thereby facilitating servicing, replacement, etc. of such components away from the confines of the vacuum chamber and perhaps reducing the chances of contamination. In one arrangement, the mere removal of the lid also removes at least one or more internal components in the same step. In other arrangements only internal components of interest are removed or perhaps even partially removed to facilitate inspection and/or maintenance while other internal components or portions of components of interest remain within the vacuum chamber. In one arrangement, the lid can be completely removed from the base of assembly **10** that may facilitate the inspection and maintenance of the internal components without the encumbrances of attachments to, or the confines of the base. In another arrangement, the external components can be removed from the lid to perhaps facilitate the inspection of the external component without substantially increasing the pressure within or contaminating the vacuum chamber. It is also contemplated that lid **14** or lid assembly **11** may remain partially coupled to base **12** in the second operable position (e.g. coupled via a hinge).

The following non-limiting examples are provided to further to facilitate aspects of the disclosure with respect to exemplary mass spectrometry operations of assembly **10**.

Methyl salicylate spectrum in FIG. **8** is obtained with internal Membrane Introduction Mass Spectrometry (MIMS)/internal Electron Ionization (EI) using a Griffin Analytical Technologies (West Lafayette, Ind.) Minotaur Model 2100A CIT Mass Spectrometer.

Perfluorodimethylcyclohexane (PDCH) spectrum of FIG. **9** is obtained with direct inlet/external glow discharge ionization using a Griffin Analytical Technologies (West Lafayette, Ind.) Minotaur Model 2100A CIT Mass Spectrometer.

Methyl salicylate spectrum of FIG. **10** is obtained with external Membrane Introduction Mass Spectrometry (MIMS)/internal Electron Ionization (EI) using a Griffin Analytical Technologies (West Lafayette, Ind.) Minotaur Model 2100A CIT Mass Spectrometer.

Dimethyl Methylphosphonate (DMMP) of FIG. **11** is obtained with Solid-Phase Microextraction (SPME)/internal Electron Ionization (EI) using a Griffin Analytical Technologies (West Lafayette, Ind.) Minotaur Model 2100A CIT Mass Spectrometer.

What is claimed is:

1. An ion trap mass spectrometer comprising:
 - a vacuum chamber housing comprising a lid and a base, the lid and the base defining a vacuum chamber volume, and the base comprising at least one wall configured to couple with the lid, wherein the lid comprises:
 - an interior surface configured to define an interior wall of the vacuum chamber and an exterior surface configured to couple with an analysis component; and
 - a first opening extending through the lid from the analysis component to the interior surface;
 - an ion trap mass separation component coupled to the interior surface of the lid without being coupled to the base and configured to perform at least some operations with respect to mass separation for use in mass spectrometry, wherein the first opening provides fluid communication between the ion trap and the analysis component;
 wherein the lid is configured to be removably operably coupled with respect to the base and positioned in a first operable position to seal with the base and provide the ion trap component wholly within the vacuum chamber volume, and a second position to decouple the lid and all analysis components coupled to the lid from the base and provide the ion trap component wholly outside the vacuum chamber volume.
2. The spectrometer of claim **1** further comprising:
 - at least one edge of the lid extending between the interior and exterior surfaces and defining a perimeter of the lid; and
 - a second opening extending from the edge through the lid to the first opening.
3. The spectrometer of claim **2** wherein the second opening is configured to receive sample for analysis.
4. The spectrometer of claim **1** wherein the analysis component comprises an ion source component configured to perform at least some operations with respect to providing ions for use in mass spectrometry.
5. The spectrometer of claim **4** further comprising a sample inlet component coupled to the interior wall of the lid and the ion trap component, the sample inlet component being operably configured to expose sample to ions received from the ion source component coupled to the exterior surface of the lid.
6. The spectrometer of claim **5** wherein the sample inlet component comprises a semi-permeable membrane.

UNITED STATES PATENT AND TRADEMARK OFFICE
CERTIFICATE OF CORRECTION

PATENT NO. : 7,427,750 B2
APPLICATION NO. : 10/542187
DATED : September 23, 2008
INVENTOR(S) : Grossenbacher et al.

Page 1 of 1

It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

Col. 5, line 41, please delete “**32-26**” after “components” and insert --**32-36**--.

Col. 7, line 30, please delete “**88**” after “surface” and insert --**86**--.

Col. 8, line 17, please delete “canter” after “the” and insert --center--.

Col. 9, line 65, please delete “to” after “ther”.

Signed and Sealed this

Twenty-fourth Day of February, 2009



JOHN DOLL
Acting Director of the United States Patent and Trademark Office